IN THE CLAIMS

The following listing of claims will replace all prior versions and listings, of claims in this application.

Claim 1 (Currently Amended): A <u>purified</u> citrullinated polypeptide <u>which reacts with</u> rheumatoid arthritis-specific autoantibodies, and is selected from the group consisting of:

- a) a citrullinated α -chain of a mammalian fibrin;
- b) a citrullinated α -chain of a mammalian fibrinogen;
- c) a fragment of at least 5 consecutive amino acids of a) and which also comprises at least one citrulline residue derived from all or part of the sequence of the α chain or of the β chain of a vertebrate fibrin, by substitution of at least one arginine residue with a citrulline residue.

Claim 2 (Cancelled).

Claim 3 (Currently Amended): The citrullinated polypeptide as claimed in claim 1, which is a) and wherein said vertebrate fibrin is a mammalian fibrin, preferably a is human fibrin.

Claim 4 (Cancelled).

Claim 5 (Currently Amended): An antigenic composition for diagnosing the presence of rheumatoid arthritis-specific autoantibodies in a biological sample, characterized in that it contains-comprising at least one citrullinated polypeptide as claimed in claim 1, optionally labeled with and/or conjugated to a carrier molecule.

Claim 6 (Currently Amended): A method for detecting rheumatoid arthritis specific autoantibodies in a biological sample, which method-is characterized in that it comprises:

bringing contacting said biological sample into contact with at least one polypeptide as claimed in claim 1, under conditions which allow the formation of an antigen/antibody complex with the rheumatoid arthritis-specific autoantibodies possibly present; and detecting, by any suitable means, the antigen/antibody complex possibly formed.

Claim 7 (Currently Amended): A kit for detecting rheumatoid arthritis-specific autoantibodies in a biological sample, characterized in that it comprises comprising at least one polypeptide as claimed in claim 1, and also buffers and reagents suitable for constituting a reaction medium which allows the formation of an antigen/antibody complex, and/or means for detecting said antigen/antibody complex.

Claims 8 and 9 (Cancelled).

Claim 10 (Currently Amended): A pharmaceutical composition, characterized in that it contains, as active principle, comprising at least one citrullinated polypeptide as claimed in claim 1, and a carrier.

Claim 11 (New): The antigenic composition according to claim 5, wherein said citrullinated polypeptide is labeled.

Claim 12 (New): The antigenic composition according to claim 5, wherein said citrullinated polypeptide is conjugated to a carrier molecule.

Claim 13 (New): The purified citrullinated polypeptide according to claim 1, which is a).

Claim 14 (New): The purified citrullinated polypeptide according to claim 1, which is b).

Claim 15 (New): The purified citrullinated polypeptide according to claim 1, which is c).

Claim 16 (New): The antigenic composition according to claim 5, wherein said citrullinated polypeptide is a).

Claim 17 (New): The antigenic composition according to claim 5, wherein said citrullinated polypeptide is b).

Claim 18 (New): The antigenic composition according to claim 5, wherein said citrullinated polypeptide is c).

Claim 19 (New): The method according to claim 6, wherein said citrullinated polypeptide is a).

Claim 20 (New): The method according to claim 6, wherein said citrullinated polypeptide is b).

Application No. 10/019,439 Reply to Office Action of August 13, 2003

Claim 21 (New): The method according to claim 6, wherein said citrullinated polypeptide is c).

Claim 22 (New): The kit according to claim 7, which further comprises reagents for detecting said antigen/antibody complex.